

## UNITED STATES OF AMERICA FEDERAL TRADE COMMISSION WASHINGTON, D.C. 20580

Bureau of Competition

May 21, 2025

Andrew Teehan General Counsel Covis Pharma GMBH 200 Connell Dr. Berkeley Heights New Jersey 07922 ateehan@covispharma.com

## *Re:* Improper Orange Book Patent Listings for Tudorza Pressair and Duaklir Pressair

Dear Mr. Teehan,

I write regarding Covis Pharma GMBH's ("Covis") ongoing obligation to ensure the propriety of its patent listings in the FDA's Approved Drug Products with Therapeutic Equivalence Evaluations (the "Orange Book"), particularly in light of the U.S. Court of Appeals for the Federal Circuit's decision in *Teva Branded Pharm. Prods. R&D, Inc. v. Amneal Pharms.* of N.Y., LLC, 124 F.4th 898 (Fed. Cir. 2024) (hereinafter "*Teva v. Amneal*").

The FTC has previously explained that patents improperly listed in the Orange Book may harm competition and delay generic drug entry, as courts have recognized.<sup>1</sup> On April 30, 2024, the FTC's Bureau of Competition (the "Bureau") sent a letter identifying a non-exhaustive list of patents that Covis had improperly submitted for listing in the Orange Book and explained how improper Orange Book listings may harm competition.<sup>2</sup> Since that letter was sent, the Federal

p239900orangebookpolicystatement092023.pdf; Brief for Fed. Trade Comm'n as Amicus Curiae, *SmithKline Beecham Corp. v. Apotex Corp.*, No. 99-CV-4304 (E.D. Pa. Jan. 28, 2003),

https://www.ftc.gov/sites/default/files/documents/amicus\_briefs/smithkline-beecham-corp.v.apotexcorp./smithklineamicus.pdf; Caraco Pharm. Labs., Ltd. v. Novo Nordisk A/S, 566 U.S. 399, 408 (2012); see also Massachusetts Laborers' Health & Welfare Fund v. Boehringer Ingelheim Pharms., Inc., No. 24-CV-10565-DJC, 2025 WL 928747, at \*20 (D. Mass. Mar. 27, 2025) ("[Plaintiff's] alleged injury, having to pay higher prices for drugs it otherwise would not need to but for [Defendants'] allegedly wrongful listing, is the precisely the kind of '[t]hreaten[ed] economic harm to consumers [that] is plainly sufficient to authorize injunctive relief.'" (quoting New York ex rel. Schneiderman v. Actavis PLC, 787 F.3d 638, 661 (2d Cir. 2015) (cleaned up)).

<sup>&</sup>lt;sup>1</sup> Fed. Trade Comm'n, Statement Concerning Brand Drug Manufacturers' Improper Listing of Patents in the Orange Book (Sept. 14, 2023), <u>https://www.ftc.gov/system/files/ftc\_gov/pdf/</u>

<sup>&</sup>lt;sup>2</sup> Apr. 30, 2024 Letter from R. Rao, Deputy Director, Bureau of Competition, to Covis Pharma GMBH, https://www.ftc.gov/system/files/ftc\_gov/pdf/covis-pharma-tudorza-and-duaklir-4302024.pdf.

Circuit's ruling in the *Teva v. Amneal* case has confirmed that the identified patents do not meet applicable Orange Book listing criteria.<sup>3</sup>

The following patents included in the Bureau's prior delisting letter remain in the Orange Book as of the date of this letter:

NDA	Product(s)	Proprietary Name	Patent Number	Listing Type
202450	1	Tudorza Pressair	8051851	DP
210595	1	Duaklir Pressair	8051851	DP

With the above patents still in the Orange Book, we are, contemporaneously with this letter, submitting patent listing dispute communications to the FDA regarding these patents. Although we have not, at this time, disputed the listing of any other Covis patents, it is Covis's responsibility to ensure that all of its patent listings comply with the statutory listing requirements, as clarified by *Teva v. Amneal*.

Combatting improper Orange Book patent listings has been a part of the FTC's longstanding enforcement and advocacy work to challenge anticompetitive conduct that stymies generic drug entry and the resulting substantial cost savings.<sup>4</sup> The FTC will remain vigilant to promote competition and protect the American public from the harms that flow from anticompetitive practices in the pharmaceutical industry.

Sincerely,

<u>/s/ Kelse Moen</u> Kelse Moen Deputy Director Bureau of Competition

<sup>4</sup> See, e.g., Biovail Corp., 134 F.T.C. 407 (2002), <u>https://www.ftc.gov/sites/default/files/documents/cases/</u> <u>2002/10/biovaildo.pdf</u>; Brief for Fed. Trade Comm'n as Amicus Curiae, Jazz Pharms., Inc. v. Avadel CNS Pharms. No. 1:21-cv-00691 (D. Del. Nov. 10, 2022), ECF No. 222-3; Brief for Fed. Trade Comm'n as Amicus Curiae, Teva Branded Pharm. Prods. R&D, Inc. v. Amneal Pharms. of N.Y., LLC, No. 24-1936 (Fed. Cir. Sept. 6, 2024), ECF No. 62; see also Mem. of Law of Amicus Curiae the Federal Trade Commission in Opp'n to Defs.' Mot. to Dismiss, In re: Buspirone Patent Litig., MDL Docket No. 1410 (S.D.N.Y. Jan. 8, 2002),

<u>https://www.ftc.gov/sites/default/files/documents/amicus\_briefs/re-buspirone-antitrust-litigation/buspirone.pdf;</u> see also Fed. Trade Comm'n, Overview of FTC Actions in Pharmaceutical Products and Distribution (Sept. 2021), <u>https://www.ftc.gov/system/files/attachments/competition-policy-guidance/overview of ftc actions in pharmaceutical products and distribution.pdf.</u>

<sup>&</sup>lt;sup>3</sup> *Teva v. Amneal*, 124 F.4th at 911 (explaining that a patent claims the drug as required for listing in the Orange Book "when it particularly points out and distinctly claims the drug as the invention.").